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FULL ESTIMATED COST

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Uploading C:\Documents and Settings\mgraffeo\My Documents\Critical Data\10533077\compound.str

chain nodes :
10 11 12
ring nodes :

1 2 3 4 5 6 7 8 9 13 14 15 16 17 18

chain bonds :

8-10 10-11 10-12 12-13

ring bonds :

 $1-2 \quad 1-6 \quad 2-3 \quad 3-4 \quad 4-5 \quad 5-6 \quad 5-7 \quad 6-9 \quad 7-8 \quad 8-9 \quad 13-14 \quad 13-18 \quad 14-15 \quad 15-16 \quad 16-17$

17-18

exact/norm bonds :

5-7 6-9 7-8 8-9 8-10 10-11 10-12

exact bonds :

12-13

normalized bonds :

1-2 1-6 2-3 3-4 4-5 5-6 13-14 13-18 14-15 15-16 16-17 17-18

Match level :

1:Atom 2:Atom 3:Atom 4:Atom 5:Atom 6:Atom 7:Atom 8:Atom 9:Atom 10:CLASS 11:CLASS 12:CLASS 13:Atom 14:Atom 15:Atom 16:Atom 17:Atom 18:Atom

L1 STRUCTURE UPLOADED

=> s l1 sss full

FULL SEARCH INITIATED 11:15:56 FILE 'REGISTRY'
FULL SCREEN SEARCH COMPLETED - 2552 TO ITERATE

100.0% PROCESSED 2552 ITERATIONS 2382 ANSWERS

SEARCH TIME: 00.00.01

L2 2382 SEA SSS FUL L1

=> file caplus

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ENTRY SESSION
FULL ESTIMATED COST 161.33 161.54

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15736 MALARIA

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OR RHINITIS OR BRONCHIECTASIS OR DERMATITIS OR MALARIA)

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bronchiectasis or dermatitis or malaria)
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         62330 ALLERG?
         28236 ASTHMA
          2027 URTICARIA
         10736 ANAPHYLAXIS
          5122 RHINITIS
           523 BRONCHIECTASIS
         15489 DERMATITIS
         15736 MALARIA
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               OR RHINITIS OR BRONCHIECTASIS OR DERMATITIS OR MALARIA)
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or bufalin or bafilomycin or concanamycin) and ige and allerg? and (asthma or
urticaria or anaphylaxis or rhinitis or bronchiectasis or dermatitis or malaria)
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       4030191 "ACID"
         52124 "OLEIC ACID"
                 ("OLEIC"(W) "ACID")
          7807 CATECHIN
            19 SCOPADULCIOL
            74 PENTAGALLOYL
           333 BUFALIN
          1157 BAFILOMYCIN
           301 CONCANAMYCIN
         19865 IGE
         62330 ALLERG?
         28236 ASTHMA
          2027 URTICARIA
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          5122 RHINITIS
           523 BRONCHIECTASIS
         15489 DERMATITIS
         15736 MALARIA
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               OR PENTAGALLOYL OR BUFALIN OR BAFILOMYCIN OR CONCANAMYCIN) AND
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               TIS OR BRONCHIECTASIS OR DERMATITIS OR MALARIA)
=> d bib abs
     ANSWER 1 OF 1 CAPLUS COPYRIGHT 2005 ACS on STN
L7
AN
     2004:412815 CAPLUS
DN
     140:386032
     Composition using a benzimidazolic compound with proton pump inhibitor
ΤI
     activity for preventing secretion of immunoglobulin E-dependent histamine
     releasing factor
     Lee, Kyung-Lim; Lee, Chul-Hee; Choi, Seung-Hee
IN
     S. Korea
PA
SO
     PCT Int. Appl., 29 pp.
     CODEN: PIXXD2
DT
     Patent
LA
     English
FAN.CNT 1
     PATENT NO.
                          KIND
                                 DATE
                                             APPLICATION NO.
                                                                     DATE
                          ----
                                 -----
                                             ------
                                            WO 2003-KR2332
                          A1
                                 20040521
                                                                     20031103
PΙ
     WO 2004041280
         W: AE, AG, AL, AM, AT, AU, AZ, BA, BB, BG, BR, BY, BZ, CA, CH, CN,
             CO, CR, CU, CZ, DE, DK, DM, DZ, EC, EE, ES, FI, GB, GD, GE, GH, GM, HR, HU, ID, IL, IN, IS, JP, KE, KG, KP, KZ, LC, LK, LR, LS,
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LT, LU, LV, MA, MD, MG, MK, MN, MW, MX, MZ, NI, NO, NZ, OM, PG,
             PH, PL, PT, RO, RU, SC, SD, SE, SG, SK, SL, SY, TJ, TM, TN, TR,
             TT, TZ, UA, UG, US, UZ, VC, VN, YU, ZA, ZM, ZW
         RW: BW, GH, GM, KE, LS, MW, MZ, SD, SL, SZ, TZ, UG, ZM, ZW, AM, AZ,
             BY, KG, KZ, MD, RU, TJ, TM, AT, BE, BG, CH, CY, CZ, DE, DK, EE,
             ES, FI, FR, GB, GR, HU, IE, IT, LU, MC, NL, PT, RO, SE, SI, SK,
             TR, BF, BJ, CF, CG, CI, CM, GA, GN, GQ, GW, ML, MR, NE, SN, TD, TG
PRAI KR 2002-67653
                          Α
                                20021102
     KR 2003-75511
                          Α
                                20031028
     The invention discloses a composition for inhibiting the secretion of an
AB
     IqE-dependent histamine-releasing factor, and pharmaceutical use
     thereof. The composition of the invention contains a benzimidazolic compound
    having proton pump inhibitor activity as an active ingredient.
     Furthermore, the composition of the invention may contain at least one
     substance selected from the group consisting of fenoctimine, oleic
     acid, catechin, scopadulciol,
    pentagalloyl glucose, bufalin, and macrolide antibiotic
    bafilomycin and concanamycin, all having proton pump
     inhibitor activity, as an addnl. active ingredient or a single active
     ingredient. The composition of the invention can be advantageously used as a
    pharmaceutical composition for the prevention and treatment of various
     allergic diseases and malaria, which are caused by the
    IgE-dependent histamine-releasing factor.
=> s 16
         19865 IGE
         62330 ALLERG?
         28236 ASTHMA
          2027 URTICARIA
         10736 ANAPHYLAXIS
          5122 RHINITIS
           523 BRONCHIECTASIS
         15489 DERMATITIS
         15736 MALARIA
             4, L3 AND IGE AND ALLERG? AND (ASTHMA OR URTICARIA OR ANAPHYLAXIS
L8
               OR RHINITIS OR BRONCHIECTASIS OR DERMATITIS OR MALARIA)
=> d 1-4 bib abs hitstr
    ANSWER 1 OF 4 CAPLUS COPYRIGHT 2005 ACS on STN
L8
    2004:834661 CAPLUS
ΑN
DN
     142:348646
TI
    Recurrent anaphylaxis linked to pantoprazole
    Kollmeier, Alexa P.; Eddleston, Jane; Zuraw, Bruce L.; Christiansen,
ΑU
    Department of Asthma, Allergy and Immunol., Scripps Clin., La Jolla, CA,
CS
     92037, USA
     Journal of Allergy and Clinical Immunology (2004), 114(4), 975-977
so
     CODEN: JACIBY; ISSN: 0091-6749
PΒ
    Elsevier Inc.
DT
     Journal
LA
     English
AΒ
     The case of a 47-yr-old man with recurrent anaphylaxis induced
    by pantoprazole, a benzimidazole proton pump inhibitor, is presented.
     this patient, the pos. skin test response and increased tryptase level are
     consistent with prior case reports of proton pump inhibitor
     anaphylaxis and suggest an immediate hypersensitivity mechanism.
    Although mutations in the CYP2C19 gene were not identified, the timing of
     anaphylactic events invokes the possible involvement of modifying
    pharmacogenetic factors, variations in relative levels of drug-specific
     IqE, or both.
     102625-70-7, Pantoprazole 138786-67-1, Protonix
IT
     RL: ADV (Adverse effect, including toxicity); THU (Therapeutic use); BIOL
```

(Biological study); USES (Uses)

(recurrent anaphylaxis linked to pantoprazole)

RN 102625-70-7 CAPLUS

CN 1H-Benzimidazole, 5-(difluoromethoxy)-2-[[(3,4-dimethoxy-2-pyridinyl)methyl]sulfinyl]- (9CI) (CA INDEX NAME)

$$F_2CH-O$$
 N
 S
 S
 CH_2
 N
 N
 N
 N
 N

RN 138786-67-1 CAPLUS

CN 1H-Benzimidazole, 5-(difluoromethoxy)-2-[[(3,4-dimethoxy-2-pyridinyl)methyl]sulfinyl]-, sodium salt (9CI) (CA INDEX NAME)

Na

RE.CNT 9 THERE ARE 9 CITED REFERENCES AVAILABLE FOR THIS RECORD ALL CITATIONS AVAILABLE IN THE RE FORMAT

L8 ANSWER 2 OF 4 CAPLUS COPYRIGHT 2005 ACS on STN

AN 2004:412815 CAPLUS

DN 140:386032

TI Composition using a benzimidazolic compound with proton pump inhibitor activity for preventing secretion of immunoglobulin E-dependent histamine releasing factor

IN Lee, Kyung-Lim; Lee, Chul-Hee; Choi, Seung-Hee

PA S. Korea

SO PCT Int. Appl., 29 pp.

CODEN: PIXXD2

DT Patent

LA English

FAN. CNT 1

FAN.	CNT	1																		
	PATENT NO.)	DATE		1	APPL	ICAT	ION I	NO.	DATE					
ΡI	WO 2004041280					A1	20040521			1	WO 2	003-	KR23:	20031103						
		W:	ΑE,	AG,	AL,	AM,	AT,	AU,	AZ,	BA,	BB,	BG,	BR,	BY,	BZ,	CA,	CH,	CN,		
			CO,	CR,	CU,	CZ,	DE,	DK,	DM,	DZ,	EC,	EE,	ES,	FI,	GB,	GD,	GE,	GH,		
			GM,	HR,	HU,	ID,	IL,	IN,	IS,	JP,	KΕ,	KG,	ΚP,	ΚZ,	LC,	LK,	LR,	LS,		
			LT,	LU,	LV,	MA,	MD,	MG,	MK,	MN,	MW,	MX,	ΜZ,	NI,	NO,	NZ,	OM,	PG,		
			PH,	PL,	PT,	RO,	RU,	SC,	SD,	SE,	SG,	SK,	SL,	SY,	ТJ,	TM,	TN,	TR,		
			TT,	TZ,	UA,	UG,	US,	UZ,	VC,	VN,	YU,	ZA,	ZM,	ZW						
		RW:	BW,	GH,	GM,	KE,	LS,	MW,	MZ,	SD,	SL,	SZ,	TZ,	UG,	ZM,	ZW,	ΑM,	AZ,		
			BY,	KG,	ΚZ,	MD,	RU,	ТJ,	TM,	AT,	BE,	BG,	CH,	CY,	CZ,	DE,	DK,	EE,		
			ES,	FI,	FR,	GB,	GR,	HU,	IE,	IT,	LU,	MC,	NL,	PT,	RO,	SE,	SI,	SK,		
			TR,	BF,	ВJ,	CF,	CG,	CI,	CM,	GΑ,	GN,	GQ,	GW,	ML,	MR,	NE,	SN,	TD,	TG	
PRAI	KR	2002	-676	53,		Α		2002	1102											
	KR 2003-75511					Α		2003	1028											
										_				_				_		

AB The invention discloses a composition for inhibiting the secretion of an IgE-dependent histamine-releasing factor, and pharmaceutical use thereof. The composition of the invention contains a benzimidazolic compound

having proton pump inhibitor activity as an active ingredient. Furthermore, the composition of the invention may contain at least one substance selected from the group consisting of fenoctimine, oleic acid, catechin, scopadulciol, pentagalloyl glucose, bufalin, and macrolide antibiotic bafilomycin and concanamycin, all having proton pump inhibitor activity, as an addnl. active ingredient or a single active ingredient. The composition of the invention can be advantageously used as a pharmaceutical composition for the prevention and treatment of various allergic diseases and malaria, which are caused by the IgE -dependent histamine-releasing factor.

73590-58-6, Omeprazole 102625-70-7, Pantoprazole
103577-45-3, Lansoprazole 117976-89-3, Rabeprazole
RL: PAC (Pharmacological activity); THU (Therapeutic use); BIOL
(Biological study); USES (Uses)

(benzimidazolic compound with proton pump inhibitor activity for preventing secretion of IgE-dependent histamine releasing factor)

RN 73590-58-6 CAPLUS

CN 1H-Benzimidazole, 5-methoxy-2-[[(4-methoxy-3,5-dimethyl-2-pyridinyl)methyl]sulfinyl]- (9CI) (CA INDEX NAME)

RN 102625-70-7 CAPLUS

CN 1H-Benzimidazole, 5-(difluoromethoxy)-2-[[(3,4-dimethoxy-2-pyridinyl)methyl]sulfinyl]- (9CI) (CA INDEX NAME)

RN 103577-45-3 CAPLUS

CN 1H-Benzimidazole, 2-[[[3-methyl-4-(2,2,2-trifluoroethoxy)-2-pyridinyl]methyl]sulfinyl]- (9CI) (CA INDEX NAME)

RN 117976-89-3 CAPLUS

CN 1H-Benzimidazole, 2-[[[4-(3-methoxypropoxy)-3-methyl-2-pyridinyl]methyl]sulfinyl]- (9CI) (CA INDEX NAME)

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L8 ANSWER 3 OF 4 CAPLUS COPYRIGHT 2005 ACS on STN
```

AN 2002:223222 CAPLUS

DN 137:257592

TI T-cell reactions to drugs in distinct clinical manifestations of drug allergy

AU Neukomm, Corinne B.; Yawalkar, Nikhil; Helbling, Arthur; Pichler, Werner J.

CS Division of Allergology, Clinic of Rheumatology and Clinical Immunology/Allergology, Inselspital, Bern, Switz.

SO Journal of Investigational Allergology and Clinical Immunology (2001), 11(4), 275-284

CODEN: JIAIEF; ISSN: 1018-9068 PB Hogrefe & Huber Publishers

DT Journal

LA English

Recent data indicate that T cells play a major role in different forms of AB drug allergies. To show that T-cell reactions are involved in various forms of adverse reactions to different kinds of drugs, and that lymphocyte transformation and skin tests may be pos. in patients who had distinct clin. manifestations of drug allergies. We collected data of 44 patients with a highly suggestive history for adverse drug reaction who had on subsequent investigations a pos. lymphocyte transformation test. In 41/44 patients (93%) skin tests with the suspected drugs were performed and in some cases drug-specific IgE -antibodies were determined All patients were HLA typed. Clin. manifestations of the drug allergy were heterogeneous, comprising maculopapular and bullous exanthema, erythema exsudativum multiforme, vasculitis, serum sickness, urticaria, as well as involvement of internal organs. Maculopapular exanthemas formed the largest group (54%), followed by reactions more indicative of immediate hypersensitivity (28%), such as urticaria/angioedema. In most cases (63%), β-lactam antibiotics were found to have caused the allergic reaction. Skin tests for immediate reactions were pos. in 6/40 patients (15%) tested, those for late reactions in 24/38 patients (63%) tested. Our data provide evidence that drug-specific T cells can be detected in distinct clin. manifestations of drug allergy. A combined approach using a detailed case history, lymphocyte transformation tests, skin tests (immediate and delayed type) appears to be helpful to identifying the incriminated drug.

IT 103577-45-3, Agopton

RL: ADV (Adverse effect, including toxicity); THU (Therapeutic use); BIOL (Biological study); USES (Uses)

(T-cell reactions to drugs in distinct clin. manifestations of drug allergy)

RN 103577-45-3 CAPLUS

CN 1H-Benzimidazole, 2-[[[3-methyl-4-(2,2,2-trifluoroethoxy)-2-pyridinyl]methyl]sulfinyl]- (9CI) (CA INDEX NAME)

RE.CNT 26 THERE ARE 26 CITED REFERENCES AVAILABLE FOR THIS RECORD ALL CITATIONS AVAILABLE IN THE RE FORMAT

ANSWER 4 OF 4 CAPLUS COPYRIGHT 2005 ACS on STN L8

AN 2002:125206 CAPLUS

DN 137:210620

TU-572, a Potent and Selective CD45 Inhibitor, Suppresses IgE ΤI -Mediated Anaphylaxis and Murine Contact Hypersensitivity

Hamaquchi, Takuya; Takahashi, Akiko; Manaka, Akira; Sato, Masakazu; Osada, AU Hiroyuki

Medicinal Research Laboratories, Taisho Pharmaceutical Co., Ltd., CS Saitama-shi, Japan

International Archives of Allergy and Immunology (2001), 126(4), 318-324 SO CODEN: IAAIEG; ISSN: 1018-2438

PΒ S. Karger AG

DT Journal

LA English

Background: CD45, receptor-type protein tyrosine phosphatases (PTPases) AB are essential components of signaling through both the T cell receptor and the B cell antigen receptor. However, the functional significance of CD45 in the signaling pathway through the high-affinity Ig (Ig) E receptor has not yet been established. In this study, we demonstrate that the potent CD45 inhibitor neg. regulates IgE-dependent anaphylaxis and contact hypersensitivity reactions. Method: We have previously found that TU-572, 2-[(4-methylthiopyridin-2-yl)methylsulfinyl]-5isopropoxybenzimidazole, had a potent and selective inhibitory effect against PTPase activity of CD45. Using a CD45 inhibitor, we examined in vitro and in vivo IgE-mediated responses. Results: TU-572 potently inhibited histamine release from rat peritoneal mast cells and mouse systemic anaphylaxis reaction using monoclonal anti-dinitrophenyl (DNP) IgE and DNP-BSA. TU-572 also suppressed the immediate-type hypersensitivity response induced by repeated epicutaneous application of trinitrochlorobenzene in BALB/c mice. Conclusion: These findings revealed that the PTPase activity of CD45 played a critical role in signal transduction of IgE-mediated anaphylaxis in vitro and in vivo. PTPase inhibitors such as TU-572 are useful in the treatment of allergic diseases.

IT 326592-39-6, TU 572

> RL: PAC (Pharmacological activity); THU (Therapeutic use); BIOL (Biological study); USES (Uses)

(TU-572, a potent and selective CD45 inhibitor, suppresses IgE -mediated anaphylaxis and murine contact hypersensitivity reactions)

326592-39-6 CAPLUS RN

1H-Benzimidazole, 5-(1-methylethoxy)-2-[[[4-(methylthio)-2-CN pyridinyl]methyl]sulfinyl]- (9CI) (CA INDEX NAME)

RE.CNT 37 THERE ARE 37 CITED REFERENCES AVAILABLE FOR THIS RECORD ALL CITATIONS AVAILABLE IN THE RE FORMAT

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LAST RELOADED: Sep 2, 2005 (20050902/UP).

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FILE COVERS 1907 - 7 Sep 2005 VOL 143 ISS 11 FILE LAST UPDATED: 6 Sep 2005 (20050906/ED)

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=> s 13 L9 4437 L2

=> sel rn
E1 THROUGH E17 ASSIGNED

=> file reg COST IN U.S. DOLLARS SINCE FILE TOTAL ENTRY SESSION FULL ESTIMATED COST 3.32 310.90 DISCOUNT AMOUNTS (FOR QUALIFYING ACCOUNTS) SINCE FILE TOTAL ENTRY SESSION CA SUBSCRIBER PRICE 0.00 -3.65

FILE 'REGISTRY' ENTERED AT 11:24:49 ON 07 SEP 2005 USE IS SUBJECT TO THE TERMS OF YOUR STN CUSTOMER AGREEMENT. PLEASE SEE "HELP USAGETERMS" FOR DETAILS. COPYRIGHT (C) 2005 American Chemical Society (ACS)

Property values tagged with IC are from the ZIC/VINITI data file provided by InfoChem.

STRUCTURE FILE UPDATES: 6 SEP 2005 HIGHEST RN 862534-94-9 DICTIONARY FILE UPDATES: 6 SEP 2005 HIGHEST RN 862534-94-9

New CAS Information Use Policies, enter HELP USAGETERMS for details.

TSCA INFORMATION NOW CURRENT THROUGH JULY 14, 2005

Please note that search-term pricing does apply when conducting SmartSELECT searches.

Structure search iteration limits have been increased. See HELP SLIMITS for details.

Experimental and calculated property data are now available. For more information enter HELP PROP at an arrow prompt in the file or refer to the file summary sheet on the web at: http://www.cas.org/ONLINE/DBSS/registryss.html

=> s e1-e17

1 102625-70-7/BI (102625-70-7/RN) 1 103577-45-3/BI (103577-45-3/RN) 1 112-80-1/BI (112-80-1/RN) 1 117976-89-3/BI (117976-89-3/RN) 1 136565-26-9/BI (136565-26-9/RN) 1 151499-39-7/BI (151499-39-7/RN) 1 154-23-4/BI (154-23-4/RN)1 4091-50-3/BI (4091-50-3/RN) 1 465-21-4/BI (465-21-4/RN) 1 50-00-0/BI (50-00-0/RN)

```
1 50-78-2/BI
                  (50-78-2/RN)
             1 50678-27-8/BI
                  (50678-27-8/RN)
             1 51-17-2/BI
                  (51-17-2/RN)
             1 51-45-6/BI
                  (51-45-6/RN)
             1 69365-65-7/BI
                  (69365-65-7/RN)
             1 73590-58-6/BI
                  (73590-58-6/RN)
             1 80890-47-7/BI
                  (80890-47-7/RN)
L11
            17 (102625-70-7/BI OR 103577-45-3/BI OR 112-80-1/BI OR 117976-89-3/
               BI OR 136565-26-9/BI OR 151499-39-7/BI OR 154-23-4/BI OR 4091-50
               -3/BI OR 465-21-4/BI OR 50-00-0/BI OR 50-78-2/BI OR 50678-27-8/B
               I OR 51-17-2/BI OR 51-45-6/BI OR 69365-65-7/BI OR 73590-58-6/BI
               OR 80890-47-7/BI)
=> d1-17
D1-17 IS NOT A RECOGNIZED COMMAND
The previous command name entered was not recognized by the system.
For a list of commands available to you in the current file, enter
"HELP COMMANDS" at an arrow prompt (=>).
=> d 1-17
L11 ANSWER 1 OF 17 REGISTRY COPYRIGHT 2005 ACS on STN
     151499-39-7 REGISTRY
RN
ED
     Entered STN: 01 Dec 1993
CN
     Bafilomycin (9CI) (CA INDEX NAME)
MF
     Unspecified
CI
     MAN
SR
     STN Files:
                  AGRICOLA, BIOBUSINESS, BIOSIS, CA, CAPLUS, TOXCENTER, USPAT2,
LC
       USPATFULL
*** STRUCTURE DIAGRAM IS NOT AVAILABLE ***
              63 REFERENCES IN FILE CA (1907 TO DATE)
               2 REFERENCES TO NON-SPECIFIC DERIVATIVES IN FILE CA
              63 REFERENCES IN FILE CAPLUS (1907 TO DATE)
L11 ANSWER 2 OF 17 REGISTRY COPYRIGHT 2005 ACS on STN
     136565-26-9 REGISTRY
RN
     Entered STN: 04 Oct 1991
ED
     9,11a-Methano-11aH-cyclohepta[a]naphthalen-8(9H)-one, 5-
CN
     (benzoyloxy) dodecahydro-4-(hydroxymethyl)-4,9,11b-trimethyl-,
     (4R, 4aR, 5R, 6aS, 9S, 11aS, 11bS) - (9CI) (CA INDEX NAME)
OTHER CA INDEX NAMES:
     9,11a-Methano-11aH-cyclohepta[a]naphthalen-8(9H)-one, 5-
     (benzoyloxy) dodecahydro-4-(hydroxymethyl)-4,9,11b-trimethyl-,
     [4R-(4\alpha, 4a\alpha, 5\beta, 6a\beta, 9\beta, 11a\beta, 11b\beta)] -
OTHER NAMES:
     Scopadulciol
CN
     C27 H36 O4
MF
SR
     CA
LC
     STN Files:
                  ADISINSIGHT, AGRICOLA, BEILSTEIN*, BIOSIS, CA, CAPLUS, IPA,
       MEDLINE, TOXCENTER
         (*File contains numerically searchable property data)
```

Me
$$CH_2-OH$$
 $O-C-Ph$
 O

PROPERTY DATA AVAILABLE IN THE 'PROP' FORMAT

13 REFERENCES IN FILE CA (1907 TO DATE)

13 REFERENCES IN FILE CAPLUS (1907 TO DATE)

L11 ANSWER 3 OF 17 REGISTRY COPYRIGHT 2005 ACS on STN

RN 117976-89-3 REGISTRY

ED Entered STN: 16 Dec 1988

CN 1H-Benzimidazole, 2-[[[4-(3-methoxypropoxy)-3-methyl-2-pyridinyl]methyl]sulfinyl]- (9CI) (CA INDEX NAME)

OTHER NAMES:

CN 2-[[[3-Methyl-4-(3-methoxypropoxy)-2-pyridyl]methyl]sulfinyl]-1Hbenzimidazole

CN 2-[[[4-(3-Methoxypropoxy)-3-methyl-2-pyridyl]methyl]sulfinyl]-1Hbenzimidazole

CN 2-[[[4-(3-Methoxypropoxy)-3-methyl-2-pyridyl]methyl]sulfinyl]benzimidazole

CN LY 307640

CN Pariets

CN Rabeprazole

FS 3D CONCORD

MF C18 H21 N3 O3 S

CI COM

SR CA

LC STN Files: ADISINSIGHT, ADISNEWS, ANABSTR, BIOSIS, BIOTECHNO, CA, CAPLUS, CASREACT, CBNB, CHEMCATS, CIN, DDFU, DIOGENES, DRUGU, EMBASE, IMSDRUGNEWS, IMSPATENTS, IMSRESEARCH, IPA, MRCK*, PATDPASPC, PHAR, PROMT, PROUSDDR, PS, SYNTHLINE, TOXCENTER, USAN, USPAT2, USPATFULL (*File contains numerically searchable property data)

Other Sources: WHO

$$\begin{array}{c|c} H & O \\ \hline N & S-CH_2 \\ \hline N & O-(CH_2)_3-OMe \end{array}$$

PROPERTY DATA AVAILABLE IN THE 'PROP' FORMAT

464 REFERENCES IN FILE CA (1907 TO DATE)

12 REFERENCES TO NON-SPECIFIC DERIVATIVES IN FILE CA

466 REFERENCES IN FILE CAPLUS (1907 TO DATE)

L11 ANSWER 4 OF 17 REGISTRY COPYRIGHT 2005 ACS on STN

RN 103577-45-3 REGISTRY

ED Entered STN: 02 Aug 1986

CN 1H-Benzimidazole, 2-[[[3-methyl-4-(2,2,2-trifluoroethoxy)-2-

```
pyridinyl]methyl]sulfinyl]- (9CI) (CA INDEX NAME)
OTHER NAMES:
     (±)-Lansoprazole
CN
     2-[[[3-Methyl-4-(2,2,2-trifluoroethoxy)-2-pyridyl]methyl]sulfinyl]-1H-
     benzimidazole
CN
     A 65006
     AG 1749
CN
     Agopton
CN
CN
     Ilsatec
     Ketian
CN
     Lancid
CN
CN
     Lanfast
CN
     Lanproton
CN
     Lansopep
CN
     Lansophed
CN
     Lansoprazole
CN
     Lansox
CN
     Lanston
     Lanz
CN
CN
     Lanzol 30
CN
     Lanzopral
CN
     Lanzor
CN
     Lapraz
CN
     Ogast
CN
     Ogastro
     PP/K-10
CN
CN
     Prevacid
CN
     Promp
CN
     Prosogan
ĊN
     Suprecid
CN
     Takepron
CN
     Ulpax
CN
     Zoton
FS
     3D CONCORD
     154727-72-7
DR
     C16 H14 F3 N3 O2 S
MF
CI
     COM
SR
     CA
                 ADISINSIGHT, ADISNEWS, AGRICOLA, ANABSTR, BEILSTEIN*,
LC
     STN Files:
       BIOBUSINESS, BIOSIS, BIOTECHNO, CA, CANCERLIT, CAPLUS, CASREACT, CBNB,
       CHEMCATS, CIN, CSCHEM, DDFU, DIOGENES, DRUGU, EMBASE, HSDB*,
       IMSDRUGNEWS, IMSPATENTS, IMSRESEARCH, IPA, MEDLINE, MRCK*, MSDS-OHS,
       PATDPASPC, PHAR, PROMT, PROUSDDR, PS, RTECS*, SCISEARCH, SYNTHLINE,
       TOXCENTER, USAN, USPAT2, USPATFULL
         (*File contains numerically searchable property data)
     Other Sources:
                      WHO
```

PROPERTY DATA AVAILABLE IN THE 'PROP' FORMAT

1246 REFERENCES IN FILE CA (1907 TO DATE)
17 REFERENCES TO NON-SPECIFIC DERIVATIVES IN FILE CA
1251 REFERENCES IN FILE CAPLUS (1907 TO DATE)

L11 ANSWER 5 OF 17 REGISTRY COPYRIGHT 2005 ACS on STN RN 102625-70-7 REGISTRY

```
ED
     Entered STN: 14 Jun 1986
CN
     1H-Benzimidazole, 5-(difluoromethoxy)-2-[[(3,4-dimethoxy-2-
     pyridinyl)methyl]sulfinyl]- (9CI) (CA INDEX NAME)
OTHER NAMES:
     5-(Difluoromethoxy)-2-[[(3,4-dimethoxy-2-pyridinyl)methyl]sulfinyl]-1H-
     benzimidazole
     5-(Difluoromethoxy)-2-[[(3,4-dimethoxy-2-pyridyl)methyl]sulfinyl]-1H-
CN
     benzimidazole
CN
     BY 1023
     Pantoprazole
CN
    Pantozol
CN
     SKF 96022
CN
     3D CONCORD
FS
DR
     154644-14-1
     C16 H15 F2 N3 O4 S
MF
CI
     COM
SR
     CA
                  ADISINSIGHT, ADISNEWS, AGRICOLA, ANABSTR, BEILSTEIN*,
LC
     STN Files:
       BIOBUSINESS, BIOSIS, BIOTECHNO, CA, CANCERLIT, CAPLUS, CASREACT, CBNB,
       CHEMCATS, CIN, CSCHEM, DDFU, DIOGENES, DRUGU, EMBASE, HSDB*,
       IMSDRUGNEWS, IMSPATENTS, IMSRESEARCH, IPA, MEDLINE, MRCK*, MSDS-OHS,
       PATDPASPC, PHAR, PROMT, PROUSDDR, RTECS*, SCISEARCH, SYNTHLINE,
       TOXCENTER, USAN, USPAT2, USPATFULL
         (*File contains numerically searchable property data)
     Other Sources:
                      WHO
```

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**PROPERTY DATA AVAILABLE IN THE 'PROP' FORMAT**
```

645 REFERENCES IN FILE CA (1907 TO DATE)

```
20 REFERENCES TO NON-SPECIFIC DERIVATIVES IN FILE CA
             649 REFERENCES IN FILE CAPLUS (1907 TO DATE)
L11 ANSWER 6 OF 17 REGISTRY COPYRIGHT 2005 ACS on STN
     80890-47-7 REGISTRY
RN
ED
     Entered STN: 16 Nov 1984
    Oxacyclooctadeca-3,5,13,15-tetraen-2-one, 18-[(1S,2R,3S)-3-[(2R,4R,5S,6R)-
CN
     4-[[4-0-(aminocarbonyl)-2,6-dideoxy-β-D-arabino-
    hexopyranosyl]oxy]tetrahydro-2-hydroxy-5-methyl-6-(1E)-1-propenyl-2H-pyran-
     2-yl]-2-hydroxy-1-methylbutyl]-9-ethyl-8,10-dihydroxy-3,17-dimethoxy-
     5,7,11,13-tetramethyl-, (3Z,5E,7R,8R,9S,10S,11R,13E,15E,17S,18R)- (9CI)
     (CA INDEX NAME)
OTHER CA INDEX NAMES:
CN
    Concanamycin A
    Oxacyclooctadecane, concanamycin A deriv.
CN
OTHER NAMES:
    Antibiotic X 4357B
CN
     Concanamycin
CN
CN
    X 4357B
     [7R-[3Z,5E,7R*,8R*,9S*,10S*,11R*,13E,15E,17S*,18R*[1S*,2R*,3S*[2R*,4R*,5S*
CN
     , 6R*(E)]]]-18-[3-[4-[[4-O-(Aminocarbonyl)-2,6-dideoxy-β-D-arabino-
     hexopyranosyl]oxy]tetrahydro-2-hydroxy-5-methyl-6-(1-propenyl)-2H-pyran-2-
     yl]-2-hydroxy-1-methylbutyl]-9-ethyl-8,10-dihydroxy-3,17-dimethoxy-
     5,7,11,13-tetramethyloxacyclooctadeca-3,5,13,15-tetraen-2-one
```

```
FS STEREOSEARCH
```

DR 66771-59-3

MF C46 H75 N O14

CI COM

LC STN Files: AGRICOLA, BIOBUSINESS, BIOSIS, BIOTECHNO, CA, CANCERLIT, CAPLUS, CASREACT, CHEMCATS, CSCHEM, DDFU, DRUGU, EMBASE, IFICDB, IFIPAT, IFIUDB, MEDLINE, NAPRALERT, RTECS*, TOXCENTER, USPAT2, USPATFULL (*File contains numerically searchable property data)

Absolute stereochemistry.

Double bond geometry as described by E or Z.

PAGE 1-B

Me

PROPERTY DATA AVAILABLE IN THE 'PROP' FORMAT

103 REFERENCES IN FILE CA (1907 TO DATE)

5 REFERENCES TO NON-SPECIFIC DERIVATIVES IN FILE CA

103 REFERENCES IN FILE CAPLUS (1907 TO DATE)

L11 ANSWER 7 OF 17 REGISTRY COPYRIGHT 2005 ACS on STN

RN 73590-58-6 REGISTRY

ED Entered STN: 16 Nov 1984

CN 1H-Benzimidazole, 5-methoxy-2-[[(4-methoxy-3,5-dimethyl-2-

pyridinyl)methyl]sulfinyl]- (9CI) (CA INDEX NAME)

OTHER NAMES:

OH

CN (\pm) -Omeprazole

CN 2-[[(3,5-Dimethyl-4-methoxy-2-pyridyl)methyl]sulfinyl]-5-methoxy-1H-benzimidazole

```
5-Methoxy-2-[[(4-methoxy-3,5-dimethyl-2-pyridyl)methyl]sulfinyl]-1H-
CN
     benzimidazole
CN
     Acidex
CN
     Antra
     Antra MUPS
CN
CN
     Audazol
CN
     Aulcer
     Belmazol
CN
CN
     Ceprandal
CN
     Desec
CN
     Dizprazol
CN
     Dudencer
CN
     Elgam
CN
     Emeproton
CN
     Epirazole
CN
     Gastrimut
CN
     GastroGard
CN
     Gastroloc
CN
     Gastrozole
CN
     Gibancer
CN
    H 168/68
CN
     Indurgan
CN
     Inhibitron
CN
     Inhipump
CN
     Logastric
     Lomac
CN
CN
     Losec
CN
     Mepral
CN
     Miol
CN
     Miracid
     Mopral
CN
     Ocid
CN
CN
     Omapren
     Omebeta 20
CN
CN
     Omed
CN
     Omedar
CN
     OMEP
CN
     Omepradex
CN
     Omepral
CN
     Omeprazen
CN
     Omeprazole
CN
     Omeprazon
CN
     Omepril
CN
     Omezol
CN
     Omezzol
CN
     Omid
CN
     Omisec
CN
     Omizac
ADDITIONAL NAMES NOT AVAILABLE IN THIS FORMAT - Use FCN, FIDE, or ALL for
     DISPLAY
     3D CONCORD
FS
     172964-80-6, 131959-78-9
DR
ΜF
     C17 H19 N3 O3 S
CI
     COM
                  ADISINSIGHT, ADISNEWS, AGRICOLA, ANABSTR, BEILSTEIN*,
LC
     STN Files:
       BIOBUSINESS, BIOSIS, BIOTECHNO, CA, CANCERLIT, CAPLUS, CASREACT, CBNB,
       CEN, CHEMCATS, CIN, CSCHEM, CSNB, DDFU, DIOGENES, DRUGU, EMBASE, HSDB*,
       IMSCOSEARCH, IMSDRUGNEWS, IMSPATENTS, IMSRESEARCH, IPA, MEDLINE, MRCK*,
       PATDPASPC, PHAR, PIRA, PROMT, PROUSDDR, PS, RTECS*, SCISEARCH,
       SYNTHLINE, TOXCENTER, USAN, USPAT2, USPATFULL, VETU
         (*File contains numerically searchable property data)
     Other Sources:
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PROPERTY DATA AVAILABLE IN THE 'PROP' FORMAT

3026 REFERENCES IN FILE CA (1907 TO DATE)
56 REFERENCES TO NON-SPECIFIC DERIVATIVES IN FILE CA
3036 REFERENCES IN FILE CAPLUS (1907 TO DATE)

L11 ANSWER 8 OF 17 REGISTRY COPYRIGHT 2005 ACS on STN

RN 69365-65-7 REGISTRY

ED Entered STN: 16 Nov 1984

CN Piperidine, 4-(diphenylmethyl)-1-[(octylimino)methyl]- (9CI) (CA INDEX NAME)

OTHER NAMES:

CN Fenoctimin

CN Fenoctimine

FS 3D CONCORD

MF C27 H38 N2

CI COM

LC STN Files: ANABSTR, BEILSTEIN*, BIOBUSINESS, BIOSIS, CA, CAPLUS, DDFU, DRUGU, EMBASE, SYNTHLINE, TOXCENTER, USAN, USPATFULL

(*File contains numerically searchable property data)

Other Sources: WHO

PROPERTY DATA AVAILABLE IN THE 'PROP' FORMAT

10 REFERENCES IN FILE CA (1907 TO DATE)

10 REFERENCES IN FILE CAPLUS (1907 TO DATE)

L11 ANSWER 9 OF 17 REGISTRY COPYRIGHT 2005 ACS on STN

RN 50678-27-8 REGISTRY

ED Entered STN: 16 Nov 1984

CN D-Glucopyranose, pentakis(3,4,5-trihydroxybenzoate) (9CI) (CA INDEX NAME)

OTHER CA INDEX NAMES:

CN D-Glucose, pentagallate (7CI)

OTHER NAMES:

CN 1,2,3,4,6-Pentagalloyl-D-glucose

CN CJ 90002

CN D-Glucose, 1,2,3,4,6-pentagallate

CN Penta-O-galloyl-D-glucose

CN Pentagalloylglucose

FS STEREOSEARCH

DR 126420-90-4, 147370-08-9, 40410-94-4

MF C41 H32 O26

LC STN Files: AGRICOLA, BEILSTEIN*, BIOBUSINESS, BIOSIS, BIOTECHNO, CA, CAOLD, CAPLUS, DDFU, DRUGU, EMBASE, IPA, MEDLINE, NAPRALERT, PHAR,

TOXCENTER, USPAT2, USPATFULL (*File contains numerically searchable property data)

Absolute stereochemistry.

PAGE 1-A

PAGE 2-A

99 REFERENCES IN FILE CA (1907 TO DATE)

OH

1 REFERENCES TO NON-SPECIFIC DERIVATIVES IN FILE CA

99 REFERENCES IN FILE CAPLUS (1907 TO DATE)

1 REFERENCES IN FILE CAOLD (PRIOR TO 1967)

```
COPYRIGHT 2005 ACS on STN
L11 ANSWER 10 OF 17 REGISTRY
RN
     4091-50-3 REGISTRY
     Entered STN: 16 Nov 1984
ED
     Benzeneethanamine, 4-methoxy-N-methyl- (9CI)
                                                     (CA INDEX NAME)
CN
OTHER CA INDEX NAMES:
     Phenethylamine, p-methoxy-N-methyl- (6CI, 8CI)
CN
OTHER NAMES:
CN
     (p-Methoxyphenethyl) methylamine
CN
     4-Methoxy-N-methylbenzeneethanamine
     4-Methoxy-N-methylphenethylamine
CN
CN
     N-(p-Methoxyphenethyl)methylamine
     N-Methyl-(p-methoxyphenethyl)amine
CN
CN
     N-Methyl-\beta-(4-methoxyphenyl)ethylamine
     N-Methyl-2-(4-methoxyphenyl)ethylamine
CN
     N-Methyl-4-methoxy-\beta-phenethylamine
CN
     N-Methyl-4-methoxyphenethylamine
CN
CN
     N-Methyl-N-(4-methoxyphenethyl)amine
```

CN p-Methoxy-N-methylphenethylamine

CN [2-(4-Methoxyphenyl)ethyl]methylamine

FS 3D CONCORD

```
MF C10 H15 N O
```

CI COM

LC STN Files: AGRICOLA, BEILSTEIN*, BIOBUSINESS, BIOSIS, CA, CANCERLIT, CAOLD, CAPLUS, CASREACT, CHEMCATS, EMBASE, IFICDB, IFIPAT, IFIUDB, IPA, MEDLINE, NAPRALERT, NIOSHTIC, RTECS*, SPECINFO, TOXCENTER, USPAT2, USPATFULL

(*File contains numerically searchable property data)

PROPERTY DATA AVAILABLE IN THE 'PROP' FORMAT

1151 REFERENCES IN FILE CA (1907 TO DATE)

745 REFERENCES TO NON-SPECIFIC DERIVATIVES IN FILE CA

1151 REFERENCES IN FILE CAPLUS (1907 TO DATE)

15 REFERENCES IN FILE CAOLD (PRIOR TO 1967)

L11 ANSWER 11 OF 17 REGISTRY COPYRIGHT 2005 ACS on STN

RN 465-21-4 REGISTRY

ED Entered STN: 16 Nov 1984

CN Bufa-20,22-dienolide, 3,14-dihydroxy-, (3β,5β)- (9CI) (CA INDEX NAME)

OTHER CA INDEX NAMES:

CN 5β -Bufa-20,22-dienolide, 3β ,14-dihydroxy- (7CI, 8CI)

CN Bufalin (6CI)

OTHER NAMES:

CN NSC 89595

FS STEREOSEARCH

DR 2381-02-4

MF C24 H34 O4

CI COM

LC STN Files: AGRICOLA, ANABSTR, BEILSTEIN*, BIOBUSINESS, BIOSIS, BIOTECHNO, CA, CANCERLIT, CAOLD, CAPLUS, CASREACT, CHEMCATS, CSCHEM, DDFU, DRUGU, EMBASE, IFICDB, IFIPAT, IFIUDB, IPA, MEDLINE, MRCK*, NAPRALERT, PROMT, RTECS*, SPECINFO, TOXCENTER, USPATFULL (*File contains numerically searchable property data)

Absolute stereochemistry.

CN

CN

NSC 2819

Sunkatol No. 1

PROPERTY DATA AVAILABLE IN THE 'PROP' FORMAT

```
296 REFERENCES IN FILE CAPLUS (1907 TO DATE)
              36 REFERENCES IN FILE CAOLD (PRIOR TO 1967)
    ANSWER 12 OF 17 REGISTRY COPYRIGHT 2005 ACS on STN
L11
RN
     154-23-4 REGISTRY
ED
     Entered STN: 16 Nov 1984
     2H-1-Benzopyran-3,5,7-triol, 2-(3,4-dihydroxyphenyl)-3,4-dihydro-,
CN
     (2R,3S) - (9CI) (CA INDEX NAME)
OTHER CA INDEX NAMES:
     2H-1-Benzopyran-3,5,7-triol, 2-(3,4-dihydroxyphenyl)-3,4-dihydro-,
CN
     (2R-trans) -
CN
     Catechol (8CI)
OTHER NAMES:
     (+) - (2R:3S) -5,7,3',4'-Tetrahydroxyflavan-3-ol
CN
     (+)-3',4',5,7-Tetrahydroxy-2,3-trans-flavan-3-ol
CN
     (+)-Catechin
CN
CN
     (+)-Catechol
     (+)-Cianidanol
CN
     (+) -Cyanidan-3-ol
CN
     (+)-Cyanidanol
CN
     (+)-Cyanidanol-3
CN
     (2R,3S)-(+)-Catechin
CN
CN
     3-Cyanidanol, (+)-
     Biocatechin
CN
     Catechin
CN
     Catechin (flavan)
CN
     Catechinic acid
CN
     Catechol (flavan)
CN
     Catechuic acid
CN
     Catergen
CN
     Cianidanol
CN
     Cyanidanol
CN
     Cyanidol
CN
CN
     D-(+)-Catechin
CN
     D-Catechin
CN
     d-Catechin
CN
     D-Catechol
CN
     Dexcyanidanol
```

295 REFERENCES IN FILE CA (1907 TO DATE)

4 REFERENCES TO NON-SPECIFIC DERIVATIVES IN FILE CA

```
Teafuran 30A
CN
CN
     trans-(+)-3,3',4',5,7-Flavanpentol
FS
     STEREOSEARCH
DR
     523994-21-0, 321-01-7, 16198-00-8, 4211-28-3, 5323-80-8, 159761-73-6,
     379227-23-3
     C15 H14 O6
MF
CI
     COM
LC
     STN Files:
                  ADISNEWS, AGRICOLA, ANABSTR, BEILSTEIN*, BIOBUSINESS, BIOSIS,
       BIOTECHNO, CA, CABA, CANCERLIT, CAOLD, CAPLUS, CASREACT, CBNB, CEN,
       CHEMCATS, CHEMINFORMRX, CHEMLIST, CIN, CSCHEM, CSNB, DDFU, DIOGENES,
       DRUGU, EMBASE, IFICDB, IFIPAT, IFIUDB, IPA, MEDLINE, MRCK*, NAPRALERT,
       NIOSHTIC, PDLCOM*, PHAR, PIRA, PROMT, PS, RTECS*, SPECINFO, TOXCENTER,
       USAN, USPAT2, USPATFULL, VETU
         (*File contains numerically searchable property data)
                     EINECS**, WHO
         (**Enter CHEMLIST File for up-to-date regulatory information)
```

Absolute stereochemistry. Rotation (+).

CN

Emersol 214NF

PROPERTY DATA AVAILABLE IN THE 'PROP' FORMAT

6273 REFERENCES IN FILE CA (1907 TO DATE)
328 REFERENCES TO NON-SPECIFIC DERIVATIVES IN FILE CA
6284 REFERENCES IN FILE CAPLUS (1907 TO DATE)
2 REFERENCES IN FILE CAOLD (PRIOR TO 1967)

L11 ANSWER 13 OF 17 REGISTRY COPYRIGHT 2005 ACS on STN RN112-80-1 REGISTRY ED Entered STN: 16 Nov 1984 9-Octadecenoic acid (9Z) - (9CI) (CA INDEX NAME) OTHER CA INDEX NAMES: CN 9-Octadecenoic acid (Z)-Oleic acid (8CI) CN OTHER NAMES: Δ9-cis-Octadecenoic acid CN Δ9-cis-Oleic acid CN CN 9-cis-Octadecenoic acid 9-Octadecenoic acid, (Z)-CN 9Z-Octadecenoic acid CN CN cis- $\Delta 9$ -Octadecenoic acid CN cis-9-Octadecenoic acid cis-Oleic acid CN CN D 100 D 100 (fatty acid) CN Edenor ATiO5 CN Edenor FTiO5 CN Emersol 205 CN CNEmersol 211 CN Emersol 213NF

```
Emersol 233
CN
CN
     Emersol 6313NF
CN
     Extra Oleic 80R
CN
     Extra Oleic 90
CN
     Extra Oleic 99
CN
     Extra Olein 80
CN
     Extra Olein 90R
CN
     Extraolein 90
CN
     Industrene 105
CN
     Lunac O-CA
CN
     Lunac O-LL
     Lunac O-P
CN
CN
     Lunac OA
CN
     NAA 35
CN
     Neo-Fat 92-04
CN
     Oleine 7503
CN
     Pamolyn 100
CN
     Priolene 6906
CN
     Priolene 6907
CN
     Priolene 6928
CN
     Priolene 6930
     Priolene 6933
CN
     Vopcolene 27
CN
     Wecoline 00
CN
CN
     Z-9-Octadecenoic acid
FS
     STEREOSEARCH
     8046-01-3, 56833-51-3, 17156-84-2
DR
MF
     C18 H34 O2
CI
     COM
LC
                  ADISNEWS, AGRICOLA, ANABSTR, AQUIRE, BEILSTEIN*, BIOBUSINESS,
     STN Files:
       BIOSIS, BIOTECHNO, CA, CABA, CANCERLIT, CAOLD, CAPLUS, CASREACT, CBNB,
       CEN, CHEMCATS, CHEMINFORMRX, CHEMLIST, CHEMSAFE, CIN, CSCHEM, CSNB,
       DDFU, DETHERM*, DIOGENES, DIPPR*, DRUGU, EMBASE, ENCOMPLIT, ENCOMPLIT2,
       ENCOMPPAT, ENCOMPPAT2, GMELIN*, HODOC*, HSDB*, IFICDB, IFIPAT, IFIUDB,
       IPA, MEDLINE, MRCK*, MSDS-OHS, NAPRALERT, NIOSHTIC, PATDPASPC, PDLCOM*,
       PIRA, PROMT, PS, RTECS*, SPECINFO, SYNTHLINE, TOXCENTER, TULSA, USAN,
       USPAT2, USPATFULL, VETU, VTB
         (*File contains numerically searchable property data)
     Other Sources: DSL**, EINECS**, TSCA**
         (**Enter CHEMLIST File for up-to-date regulatory information)
```

Double bond geometry as shown.

$$_{\text{HO}_2\text{C}}$$
 (CH₂) 7 $_{\text{Z}}$ (CH₂) 7

PROPERTY DATA AVAILABLE IN THE 'PROP' FORMAT

45296 REFERENCES IN FILE CA (1907 TO DATE)
2559 REFERENCES TO NON-SPECIFIC DERIVATIVES IN FILE CA
45372 REFERENCES IN FILE CAPLUS (1907 TO DATE)
11 REFERENCES IN FILE CAOLD (PRIOR TO 1967)

L11 ANSWER 14 OF 17 REGISTRY COPYRIGHT 2005 ACS on STN

RN 51-45-6 REGISTRY

ED Entered STN: 16 Nov 1984

CN 1H-Imidazole-4-ethanamine (9CI) (CA INDEX NAME)

OTHER CA INDEX NAMES:

CN Histamine (8CI)

OTHER NAMES:

CN β-Imidazolyl-4-ethylamine

CN 2-(1H-Imidazol-4-yl)ethanamine

```
2-(1H-Imidazol-4-yl)ethylamine
CN
     2-(1H-Imidazol-5-yl)ethanamine
CN
CN
     2-(1H-Imidazol-5-yl)ethylamine
     2-(4-Imidazolyl)ethanamine
CN
CN
     2-(4-Imidazolyl)ethylamine
CN
     4-(2-Aminoethyl)imidazole
CN
     5-Imidazoleethylamine
CN
     Eramin
     Ergamine
CN
     Ergotidine
CN
     Imidazole-4-ethylamine
CN
CN
     NSC 33792
FS
     3D CONCORD
MF
     C5 H9 N3
CI
     COM
LC
     STN Files:
                ADISINSIGHT, ADISNEWS, AGRICOLA, ANABSTR, BEILSTEIN*,
       BIOBUSINESS, BIOSIS, BIOTECHNO, CA, CABA, CANCERLIT, CAOLD, CAPLUS,
       CASREACT, CBNB, CEN, CHEMCATS, CHEMINFORMRX, CHEMLIST, CIN, CSCHEM,
       CSNB, DDFU, DIOGENES, DRUGU, EMBASE, GMELIN*, HODOC*, HSDB*, IFICDB,
       IFIPAT, IFIUDB, IMSPATENTS, IMSRESEARCH, IPA, MEDLINE, MRCK*, MSDS-OHS,
       NAPRALERT, NIOSHTIC, PIRA, PROMT, RTECS*, SCISEARCH, SPECINFO,
       SYNTHLINE, TOXCENTER, USAN, USPAT2, USPATFULL, VETU
         (*File contains numerically searchable property data)
     Other Sources: DSL**, EINECS**, TSCA**
         (**Enter CHEMLIST File for up-to-date regulatory information)
       CH2-CH2-NH2
**PROPERTY DATA AVAILABLE IN THE 'PROP' FORMAT**
           34678 REFERENCES IN FILE CA (1907 TO DATE)
             479 REFERENCES TO NON-SPECIFIC DERIVATIVES IN FILE CA
           34687 REFERENCES IN FILE CAPLUS (1907 TO DATE)
              10 REFERENCES IN FILE CAOLD (PRIOR TO 1967)
L11 ANSWER 15 OF 17 REGISTRY COPYRIGHT 2005 ACS on STN
     51-17-2 REGISTRY
RN
ED
     Entered STN: 16 Nov 1984
     1H-Benzimidazole (9CI) (CA INDEX NAME)
OTHER CA INDEX NAMES:
CN Benzimidazole (6CI, 8CI)
OTHER NAMES:
    1,3-Benzodiazole
CN
     1,3-Diazaindene
CN
     3-Azaindole
CN
     Azindole
CN
     Benziminazole
CN
     Benzoglyoxaline
CN
     Benzoimidazole
CN
     BZI
     N, N'-Methenyl-o-phenylenediamine
CN
     NSC 759
CN
     o-Benzimidazole
CN
FS
     3D CONCORD
     25463-25-6, 79351-71-6, 116421-27-3
DR
     C7 H6 N2
MF
CI
     COM, RPS
                  ADISNEWS, AGRICOLA, ANABSTR, AQUIRE, BEILSTEIN*, BIOBUSINESS,
LC
     STN Files:
       BIOSIS, BIOTECHNO, CA, CABA, CANCERLIT, CAOLD, CAPLUS, CASREACT, CBNB,
```

CEN, CHEMCATS, CHEMINFORMRX, CHEMLIST, CIN, CSCHEM, DDFU, DETHERM*, DRUGU, EMBASE, GMELIN*, HODOC*, HSDB*, IFICDB, IFIPAT, IFIUDB, IPA, MEDLINE, MRCK*, MSDS-OHS, NAPRALERT, NIOSHTIC, PIRA, PROMT, RTECS*, SPECINFO, SYNTHLINE, TOXCENTER, ULIDAT, USPAT2, USPATFULL, VETU, VTB (*File contains numerically searchable property data)

Other Sources: EINECS**, NDSL**, TSCA**

(**Enter CHEMLIST File for up-to-date regulatory information)

CN

CN

CN

Asatard Ascoden 30

Ascolong

Ascriptin

PROPERTY DATA AVAILABLE IN THE 'PROP' FORMAT

6040 REFERENCES IN FILE CA (1907 TO DATE)
1881 REFERENCES TO NON-SPECIFIC DERIVATIVES IN FILE CA
6047 REFERENCES IN FILE CAPLUS (1907 TO DATE)
11 REFERENCES IN FILE CAOLD (PRIOR TO 1967)

L11 ANSWER 16 OF 17 REGISTRY COPYRIGHT 2005 ACS on STN RN 50-78-2 REGISTRY Entered STN: 16 Nov 1984 ED Benzoic acid, 2-(acetyloxy) - (9CI) (CA INDEX NAME) CNOTHER NAMES: 2-(Acetyloxy)benzoic acid CN2-Acetoxybenzoic acid CN2-Carboxyphenyl acetate CNCNA.S.A. Empirin CNAC 5230 CNAcenterine CNAcesal CN Acesan CNAcetard CNAceticyl CNAcetilum acidulatum CNAcetisal CNAcetol CNAcetonyl CNAcetophen CNAcetosal Acetosalic acid CNCNAcetosalin CNAcetylin Acetylsal CNCNAcetylsalicylic acid CNAcetyonyl CNAcetysal Acidum acetylsalicylicum CNÇN Acimetten CN Acisal CN Acylpyrin CN Adiro CN Albyl E CN ASA CN Asaflow CNAsagran

```
Aspalon
CN
     Aspergum
CN
CN
     Aspirdrops
CN
     Aspirin
CN
     Aspirin Protect 100
CN
     Aspirin Protect 300
CN
     Aspirin-Direkt
CN
     Aspirina 03
CN
     Aspro
CN
     Aspro Clear
CN
    Aspropharm
CN
    Asteric
CN
     Bayer
CN
     Benaspir
ADDITIONAL NAMES NOT AVAILABLE IN THIS FORMAT - Use FCN, FIDE, or ALL for
     DISPLAY
FS
     3D CONCORD
     11126-35-5, 11126-37-7, 98201-60-6, 2349-94-2, 26914-13-6
DR
MF
     C9 H8 O4
CI
     COM
LC
     STN Files:
                  ADISNEWS, AGRICOLA, ANABSTR, AQUIRE, BEILSTEIN*, BIOBUSINESS,
       BIOSIS, BIOTECHNO, CA, CABA, CANCERLIT, CAOLD, CAPLUS, CASREACT, CBNB,
       CEN, CHEMCATS, CHEMINFORMRX, CHEMLIST, CIN, CSCHEM, CSNB, DDFU,
       DETHERM*, DIOGENES, DIPPR*, DRUGU, EMBASE, GMELIN*, HODOC*, HSDB*,
       IFICDB, IFIPAT, IFIUDB, IMSCOSEARCH, IPA, MEDLINE, MRCK*, MSDS-OHS,
       NAPRALERT, NIOSHTIC, PATDPASPC, PDLCOM*, PHAR, PIRA, PROMT, PROUSDDR,
       PS, RTECS*, SCISEARCH, SPECINFO, SYNTHLINE, TOXCENTER, TULSA, ULIDAT,
       USAN, USPATZ, USPATFULL, VETU, VTB
         (*File contains numerically searchable property data)
     Other Sources: DSL**, EINECS**, TSCA**
         (**Enter CHEMLIST File for up-to-date regulatory information)
```

PROPERTY DATA AVAILABLE IN THE 'PROP' FORMAT

19146 REFERENCES IN FILE CA (1907 TO DATE)
372 REFERENCES TO NON-SPECIFIC DERIVATIVES IN FILE CA
19191 REFERENCES IN FILE CAPLUS (1907 TO DATE)
1 REFERENCES IN FILE CAOLD (PRIOR TO 1967)

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L11 ANSWER 17 OF 17 REGISTRY COPYRIGHT 2005 ACS on STN
RN
     50-00-0 REGISTRY
ED
     Entered STN: 16 Nov 1984
     Formaldehyde (8CI, 9CI) (CA INDEX NAME)
OTHER NAMES:
CN
    BFV
     F-gen
CN
CN
     Fannoform
CN
     Floquard 1015
     FM 282
CN
     Fordor
CN
     Formalin
CN
CN
     Formalith
CN
     Formic aldehyde
CN
     Formol
CN
     Fyde
CN
     Lysoform
CN
     Methaldehyde
```

```
CN
     Methanal
CN
     Methyl aldehyde
CN
     Methylene oxide
CN
     Morbicid
CN
     NSC 298885
CN
     Oxomethane
CN
     Oxymethylene
CN
     Paraform
CN
     Superlysoform
FS
     3D CONCORD
     8005-38-7, 8006-07-3, 8013-13-6, 112068-71-0
DR
MF
     C H2 O
CI
     COM
LC
     STN Files:
                  ADISNEWS, AGRICOLA, ANABSTR, AQUIRE, BEILSTEIN*, BIOBUSINESS,
       BIOSIS, BIOTECHNO, CA, CABA, CANCERLIT, CAOLD, CAPLUS, CASREACT, CBNB,
       CEN, CHEMCATS, CHEMINFORMRX, CHEMLIST, CHEMSAFE, CIN, CSCHEM, CSNB,
       DDFU, DETHERM*, DIOGENES, DIPPR*, DRUGU, EMBASE, ENCOMPLIT, ENCOMPLIT2,
       ENCOMPPAT, ENCOMPPAT2, GMELIN*, HODOC*, HSDB*, IFICDB, IFIPAT, IFIUDB,
       IPA, MEDLINE, MRCK*, MSDS-OHS, NAPRALERT, NIOSHTIC, PDLCOM*, PIRA,
       PROMT, PS, RTECS*, SCISEARCH, SPECINFO, TOXCENTER, TULSA, ULIDAT, USAN,
       USPAT2, USPATFULL, VETU, VTB
         (*File contains numerically searchable property data)
     Other Sources: DSL**, EINECS**, TSCA**
         (**Enter CHEMLIST File for up-to-date regulatory information)
```

 $H_2C = 0$

PROPERTY DATA AVAILABLE IN THE 'PROP' FORMAT

68262 REFERENCES IN FILE CA (1907 TO DATE)
6511 REFERENCES TO NON-SPECIFIC DERIVATIVES IN FILE CA
68334 REFERENCES IN FILE CAPLUS (1907 TO DATE)
19 REFERENCES IN FILE CAOLD (PRIOR TO 1967)

=> file caplus						
COST IN U.S. DOLLARS	SINCE FILE	TOTAL				
	ENTRY	SESSION				
FULL ESTIMATED COST	33.43	344.33				
DISCOUNT AMOUNTS (FOR QUALIFYING ACCOUNTS)	SINCE FILE	TOTAL				
	ENTRY	SESSION				
CA SUBSCRIBER PRICE	0.00	-3.65				

FILE 'CAPLUS' ENTERED AT 11:27:41 ON 07 SEP 2005 USE IS SUBJECT TO THE TERMS OF YOUR STN CUSTOMER AGREEMENT. PLEASE SEE "HELP USAGETERMS" FOR DETAILS. COPYRIGHT (C) 2005 AMERICAN CHEMICAL SOCIETY (ACS)

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FILE COVERS 1907 - 7 Sep 2005 VOL 143 ISS 11 FILE LAST UPDATED: 6 Sep 2005 (20050906/ED)

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New CAS Information Use Policies, enter HELP USAGETERMS for details.
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This file contains CAS Registry Numbers for easy and accurate substance identification.

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=> s 117976-89-3/rn
             466 117976-89-3
              12 117976-89-3D
L12
             457 117976-89-3/RN
                     (117976-89-3 (NOTL) 117976-89-3D)
=> s 117976-89-3/rn or 103577-45-3/rn or 154644-14-1/rn or 172964-80-6/rn or
131959-78-9/rn
             466 117976-89-3
              12 117976-89-3D
             457 117976-89-3/RN
                     (117976-89-3 (NOTL) 117976-89-3D)
            1251 103577-45-3
              17 103577-45-3D
            1241 103577-45-3/RN
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               0 154644-14-1
               0 154644-14-1D
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               0 172964-80-6D
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               0 131959-78-9D
               0 131959-78-9/RN
                     (131959-78-9 (NOTL) 131959-78-9D )
            1434 117976-89-3/RN OR 103577-45-3/RN OR 154644-14-1/RN OR 172964-80-
L13
                  6/RN OR 131959-78-9/RN
=> s 113 and 17
               1 L13 AND L7
L14
=> d bib abs hitstr
     ANSWER 1 OF 1 CAPLUS COPYRIGHT 2005 ACS on STN
AN
      2004:412815 CAPLUS
DN
      140:386032
TТ
      Composition using a benzimidazolic compound with proton pump inhibitor
      activity for preventing secretion of immunoglobulin E-dependent histamine
      releasing factor
      Lee, Kyung-Lim; Lee, Chul-Hee; Choi, Seung-Hee
IN
PA
      S. Korea
SO
      PCT Int. Appl., 29 pp.
      CODEN: PIXXD2
DT
      Patent
LA
     English
FAN.CNT 1
      PATENT NO.
                              KIND
                                       DATE
                                                      APPLICATION NO.
                                                                                  DATE
                              ____
                                                      _______
                                                      WO 2003-KR2332
ΡI
      WO 2004041280
                               A1
                                       20040521
                                                                                  20031103
          W: AE, AG, AL, AM, AT, AU, AZ, BA, BB, BG, BR, BY, BZ, CA, CH, CN, CO, CR, CU, CZ, DE, DK, DM, DZ, EC, EE, ES, FI, GB, GD, GE, GH, GM, HR, HU, ID, IL, IN, IS, JP, KE, KG, KP, KZ, LC, LK, LR, LS, LT, LU, LV, MA, MD, MG, MK, MN, MW, MX, MZ, NI, NO, NZ, OM, PG,
          PH, PL, PT, RO, RU, SC, SD, SE, SG, SK, SL, SY, TJ, TM, TN, TR, TT, TZ, UA, UG, US, UZ, VC, VN, YU, ZA, ZM, ZW
RW: BW, GH, GM, KE, LS, MW, MZ, SD, SL, SZ, TZ, UG, ZM, ZW, AM, AZ,
```

BY, KG, KZ, MD, RU, TJ, TM, AT, BE, BG, CH, CY, CZ, DE, DK, EE, ES, FI, FR, GB, GR, HU, IE, IT, LU, MC, NL, PT, RO, SE, SI, SK, TR, BF, BJ, CF, CG, CI, CM, GA, GN, GQ, GW, ML, MR, NE, SN, TD, TG PRAI KR 2002-67653 Α 20021102 KR 2003-75511 Α 20031028 AB The invention discloses a composition for inhibiting the secretion of an IgE-dependent histamine-releasing factor, and pharmaceutical use thereof. The composition of the invention contains a benzimidazolic compound having proton pump inhibitor activity as an active ingredient. Furthermore, the composition of the invention may contain at least one substance selected from the group consisting of fenoctimine, oleic acid, catechin, scopadulciol, pentagalloyl glucose, bufalin, and macrolide antibiotic bafilomycin and concanamycin, all having proton pump inhibitor activity, as an addnl. active ingredient or a single active ingredient. The composition of the invention can be advantageously used as a pharmaceutical composition for the prevention and treatment of various allergic diseases and malaria, which are caused by the IgE-dependent histamine-releasing factor. IT 73590-58-6, Omeprazole 102625-70-7, Pantoprazole 103577-45-3, Lansoprazole 117976-89-3, Rabeprazole RL: PAC (Pharmacological activity); THU (Therapeutic use); BIOL (Biological study); USES (Uses) (benzimidazolic compound with proton pump inhibitor activity for preventing secretion of IgE-dependent histamine releasing

factor)

73590-58-6 CAPLUS

RN

CN

RN 102625-70-7 CAPLUS
CN 1H-Benzimidazole, 5-(difluoromethoxy)-2-[[(3,4-dimethoxy-2-pyridinyl)methyl]sulfinyl]- (9CI) (CA INDEX NAME)

1H-Benzimidazole, 5-methoxy-2-[[(4-methoxy-3,5-dimethyl-2-

pyridinyl)methyl]sulfinyl]- (9CI) (CA INDEX NAME)

$$F_2CH-O$$
 N
 S
 CH_2
 N
 N
 N
 N
 N
 N
 N
 N

RN 103577-45-3 CAPLUS
CN 1H-Benzimidazole, 2-[[[3-methyl-4-(2,2,2-trifluoroethoxy)-2pyridinyl]methyl]sulfinyl]- (9CI) (CA INDEX NAME)

117976-89-3 CAPLUS RN 1H-Benzimidazole, 2-[[[4-(3-methoxypropoxy)-3-methyl-2-CN pyridinyl]methyl]sulfinyl]- (9CI) (CA INDEX NAME)

$$\begin{array}{c|c}
H & O \\
N & S - CH_2 \\
N & O - (CH_2)_3 - OMe
\end{array}$$

=> s l13 adn l8 MISSING OPERATOR L13 ADN The search profile that was entered contains terms or nested terms that are not separated by a logical operator.

=> s 113 and 18 2 L13 AND L8

=> d 1-2 bib abs hitstr

L15 ANSWER 1 OF 2 CAPLUS COPYRIGHT 2005 ACS on STN

2004:412815 CAPLUS AN

DN 140:386032

ΤI Composition using a benzimidazolic compound with proton pump inhibitor activity for preventing secretion of immunoglobulin E-dependent histamine releasing factor

Lee, Kyung-Lim; Lee, Chul-Hee; Choi, Seung-Hee IN

PA S. Korea

SO PCT Int. Appl., 29 pp.

CODEN: PIXXD2

DTPatent

English LA

EAM CMT 1

FAN.	TM.T.	T																		
	PAT	CENT 1	NO.			KIN	D 1	DATE		7	APPL	ICAT:	ION I		DATE					
PI	WO 2004041280					A1		20040521		Ţ	WO 2	003-1	KR23		20031103					
		W:	ΑE,	AG,	AL,	AM,	AT,	AU,	AZ,	BA,	BB,	BG,	BR,	BY,	ΒZ,	CA,	CH,	CN,		
			CO,	CR,	CU,	CZ,	DE,	DK,	DM,	DZ,	EC,	EE,	ES,	FI,	GB,	GD,	GΕ,	GH,		
			GM,	HR,	HU,	ID,	IL,	IN,	IS,	JΡ,	KΕ,	KG,	ΚP,	KZ,	LC,	LK,	LR,	LS,		
			LT,	LU,	LV,	MA,	MD,	MG,	MK,	MN,	MW,	MX,	MZ,	NI,	NO,	NZ,	OM,	PG,		
			PH,	PL,	PT,	RO,	RU,	SC,	SD,	SE,	SG,	SK,	SL,	SY,	TJ,	TM,	TN,	TR,		
			TT,	ŤΖ,	UA,	UG,	US,	UΖ,	VC,	VN,	YU,	ZA,	ZM,	zw						
		RW:	BW,	GH,	GM,	KΕ,	LS,	MW,	MZ,	SD,	SL,	SZ,	TZ,	UG,	ZM,	ZW,	AM,	ΑZ,		
			ΒY,	KG,	ΚZ,	MD,	RU,	TJ,	TM,	ΑT,	BE,	BG,	CH,	CY,	CZ,	DE,	DK,	EE,		
			ES,	FI,	FR,	GB,	GR,	HU,	ΙE,	IT,	LU,	MC,	NL,	PT,	RO,	SE,	SI,	SK,		
	•		TR,	BF,	В J ,	CF,	CG,	CI,	CM,	GA,	GN,	GQ,	GW,	ML,	MR,	ΝE,	SN,	TD,	TG	
PRAI	I KR 2002-67653					A		2002	1102											
						7		2003	1028											

20031028 KR 2003-75511

AB The invention discloses a composition for inhibiting the secretion of an IgE-dependent histamine-releasing factor, and pharmaceutical use thereof. The composition of the invention contains a benzimidazolic compound having proton pump inhibitor activity as an active ingredient. Furthermore, the composition of the invention may contain at least one substance selected from the group consisting of fenoctimine, oleic acid, catechin, scopadulciol, pentagalloyl glucose, bufalin, and macrolide antibiotic bafilomycin and concanamycin, all having proton pump inhibitor activity, as an addnl. active ingredient or a single active ingredient. The composition of the invention can be advantageously used as a pharmaceutical composition for the prevention and treatment of various allergic

diseases and malaria, which are caused by the IgE -dependent histamine-releasing factor.

IT 73590-58-6, Omeprazole 102625-70-7, Pantoprazole 103577-45-3, Lansoprazole 117976-89-3, Rabeprazole

RL: PAC (Pharmacological activity); THU (Therapeutic use); BIOL (Biological study); USES (Uses)

(benzimidazolic compound with proton pump inhibitor activity for preventing secretion of IgE-dependent histamine releasing factor)

RN 73590-58-6 CAPLUS

CN 1H-Benzimidazole, 5-methoxy-2-[[(4-methoxy-3,5-dimethyl-2-pyridinyl)methyl]sulfinyl]- (9CI) (CA INDEX NAME)

RN 102625-70-7 CAPLUS

CN 1H-Benzimidazole, 5-(difluoromethoxy)-2-[[(3,4-dimethoxy-2-pyridinyl)methyl]sulfinyl]- (9CI) (CA INDEX NAME)

RN 103577-45-3 CAPLUS

CN 1H-Benzimidazole, 2-[[[3-methyl-4-(2,2,2-trifluoroethoxy)-2-pyridinyl]methyl]sulfinyl]- (9CI) (CA INDEX NAME)

RN 117976-89-3 CAPLUS

CN 1H-Benzimidazole, 2-[[[4-(3-methoxypropoxy)-3-methyl-2-pyridinyl]methyl]sulfinyl]- (9CI) (CA INDEX NAME)

$$\begin{array}{c|c} H & O \\ \hline N & S - CH_2 \\ \hline N & O - (CH_2)_3 - OMe \end{array}$$

L15 ANSWER 2 OF 2 CAPLUS COPYRIGHT 2005 ACS on STN AN 2002:223222 CAPLUS

DN 137:257592

TI T-cell reactions to drugs in distinct clinical manifestations of drug allergy

AU Neukomm, Corinne B.; Yawalkar, Nikhil; Helbling, Arthur; Pichler, Werner J.

CS Division of Allergology, Clinic of Rheumatology and Clinical Immunology/Allergology, Inselspital, Bern, Switz.

SO Journal of Investigational Allergology and Clinical Immunology (2001), 11(4), 275-284 CODEN: JIAIEF; ISSN: 1018-9068

Hogrefe & Huber Publishers

PB Hogrefe & FDT Journal

LA English

AB Recent data indicate that T cells play a major role in different forms of drug allergies. To show that T-cell reactions are involved in various forms of adverse reactions to different kinds of drugs, and that lymphocyte transformation and skin tests may be pos. in patients who had distinct clin. manifestations of drug allergies. We collected data of 44 patients with a highly suggestive history for adverse drug reaction who had on subsequent investigations a pos. lymphocyte transformation test. In 41/44 patients (93%) skin tests with the suspected drugs were performed and in some cases drug-specific IgE -antibodies were determined All patients were HLA typed. Clin. manifestations of the drug allergy were heterogeneous, comprising maculopapular and bullous exanthema, erythema exsudativum multiforme, vasculitis, serum sickness, urticaria, as well as involvement of internal organs. Maculopapular exanthemas formed the largest group (54%), followed by reactions more indicative of immediate hypersensitivity (28%), such as urticaria/angioedema. In most cases (63%), β-lactam antibiotics were found to have caused the allergic reaction. Skin tests for immediate reactions were pos. in 6/40 patients (15%) tested, those for late reactions in 24/38 patients (63%) tested. Our data provide evidence that drug-specific T cells can be detected in distinct clin. manifestations of drug allergy. A combined approach using a detailed case history, lymphocyte transformation tests, skin tests (immediate and delayed type) appears to be helpful to identifying the incriminated drug.

IT 103577-45-3, Agopton

RL: ADV (Adverse effect, including toxicity); THU (Therapeutic use); BIOL (Biological study); USES (Uses)

(T-cell reactions to drugs in distinct clin. manifestations of drug allergy)

RN 103577-45-3 CAPLUS

CN 1H-Benzimidazole, 2-[[[3-methyl-4-(2,2,2-trifluoroethoxy)-2-pyridinyl]methyl]sulfinyl]- (9CI) (CA INDEX NAME)

$$\begin{array}{c|c}
N & \text{Me} \\
S - CH_2 & \text{NH}
\end{array}$$

$$\begin{array}{c|c}
O - CH_2 - CF_3 \\
N & \text{NH}$$

RE.CNT 26 THERE ARE 26 CITED REFERENCES AVAILABLE FOR THIS RECORD ALL CITATIONS AVAILABLE IN THE RE FORMAT

=> s (113 or 13) and 17 L16 1 (L13 OR L3) AND L7

=> d bib

L16 ANSWER 1 OF 1 CAPLUS COPYRIGHT 2005 ACS on STN

```
2004:412815 CAPLUS
AN
DN
     140:386032
     Composition using a benzimidazolic compound with proton pump inhibitor
ΤI
     activity for preventing secretion of immunoglobulin E-dependent histamine
     releasing factor
IN
     Lee, Kyung-Lim; Lee, Chul-Hee; Choi, Seung-Hee
     S. Korea
PA
     PCT Int. Appl., 29 pp.
SO
     CODEN: PIXXD2
DT
     Patent
     English
LA
FAN.CNT 1
     PATENT NO.
                        KIND
                                          APPLICATION NO.
                               DATE
     -----
                         ----
                                -----
                                           ______
                                                                  -----
                               20040521
                                          WO 2003-KR2332
ΡI
     WO 2004041280
                         A1
                                                                  20031103
            AE, AG, AL, AM, AT, AU, AZ, BA, BB, BG, BR, BY, BZ, CA, CH, CN,
             CO, CR, CU, CZ, DE, DK, DM, DZ, EC, EE, ES, FI, GB, GD, GE, GH,
             GM, HR, HU, ID, IL, IN, IS, JP, KE, KG, KP, KZ, LC, LK, LR, LS,
             LT, LU, LV, MA, MD, MG, MK, MN, MW, MX, MZ, NI, NO, NZ, OM, PG,
             PH, PL, PT, RO, RU, SC, SD, SE, SG, SK, SL, SY, TJ, TM, TN, TR,
             TT, TZ, UA, UG, US, UZ, VC, VN, YU, ZA, ZM, ZW
         RW: BW, GH, GM, KE, LS, MW, MZ, SD, SL, SZ, TZ, UG, ZM, ZW, AM, AZ,
             BY, KG, KZ, MD, RU, TJ, TM, AT, BE, BG, CH, CY, CZ, DE, DK, EE,
             ES, FI, FR, GB, GR, HU, IE, IT, LU, MC, NL, PT, RO, SE, SI, SK,
             TR, BF, BJ, CF, CG, CI, CM, GA, GN, GQ, GW, ML, MR, NE, SN, TD, TG
PRAI KR 2002-67653
                         Α
                               20021102
     KR 2003-75511
                         Α
                                20031028
=> s (113 or 13) and 18
L17
             4 (L13 OR L3) AND L8
=> d 1-4 bib abs hitstr
     ANSWER 1 OF 4 CAPLUS COPYRIGHT 2005 ACS on STN
L17
AN
     2004:834661 CAPLUS
DN
     142:348646
TI
     Recurrent anaphylaxis linked to pantoprazole
     Kollmeier, Alexa P.; Eddleston, Jane; Zuraw, Bruce L.; Christiansen,
AU
     Department of Asthma, Allergy and Immunol., Scripps Clin., La Jolla, CA,
CS
     92037, USA
SO
     Journal of Allergy and Clinical Immunology (2004), 114(4), 975-977
     CODEN: JACIBY; ISSN: 0091-6749
PB
     Elsevier Inc.
     Journal
DT
LA
     English
AB
     The case of a 47-yr-old man with recurrent anaphylaxis induced
     by pantoprazole, a benzimidazole proton pump inhibitor, is presented.
     this patient, the pos. skin test response and increased tryptase level are
     consistent with prior case reports of proton pump inhibitor
     anaphylaxis and suggest an immediate hypersensitivity mechanism.
     Although mutations in the CYP2C19 gene were not identified, the timing of
     anaphylactic events invokes the possible involvement of modifying
     pharmacogenetic factors, variations in relative levels of drug-specific
     IgE, or both.
TΤ
     102625-70-7, Pantoprazole 138786-67-1, Protonix
     RL: ADV (Adverse effect, including toxicity); THU (Therapeutic use); BIOL
     (Biological study); USES (Uses)
        (recurrent anaphylaxis linked to pantoprazole)
RN
     102625-70-7 CAPLUS
     1H-Benzimidazole, 5-(difluoromethoxy)-2-[[(3,4-dimethoxy-2-
CN
     pyridinyl)methyl]sulfinyl]- (9CI) (CA INDEX NAME)
```

RN 138786-67-1 CAPLUS

CN 1H-Benzimidazole, 5-(difluoromethoxy)-2-[[(3,4-dimethoxy-2-pyridinyl)methyl]sulfinyl]-, sodium salt (9CI) (CA INDEX NAME)

Na

RE.CNT 9 THERE ARE 9 CITED REFERENCES AVAILABLE FOR THIS RECORD ALL CITATIONS AVAILABLE IN THE RE FORMAT

L17 ANSWER 2 OF 4 CAPLUS COPYRIGHT 2005 ACS on STN

AN 2004:412815 CAPLUS

DN 140:386032

TI Composition using a benzimidazolic compound with proton pump inhibitor activity for preventing secretion of immunoglobulin E-dependent histamine releasing factor

IN Lee, Kyung-Lim; Lee, Chul-Hee; Choi, Seung-Hee

PA S. Korea

SO PCT Int. Appl., 29 pp.

CODEN: PIXXD2

DT Patent

LA English

FAN. CNT 1

FAN.	CN.L.	1																			
	PATENT NO.)	DATE		i	APPL:	ICAT	ION I		DATE						
ΡI	WO	WO 2004041280				A1	20040521			WO 2003-KR2332						20031103					
		W:	ΑE,	AG,	AL,	AM,	AT,	AU,	ΑZ,	BA,	BB,	BG,	BR,	BY,	ΒZ,	CA,	CH,	CN,			
			CO,	CR,	CU,	CZ,	DE,	DK,	DM,	DZ,	EC,	EE,	ES,	FΊ,	GB,	GD,	GE,	GH,			
			GM,	HR,	HU,	ID,	IL,	IN,	IS,	JP,	KE,	KG,	ΚP,	KΖ,	LC,	LK,	LR,	LS,			
			LT,	LU,	LV,	MA,	MD,	MG,	MK,	MN,	MW,	MX,	MZ,	NI,	NO,	NZ,	OM,	PG,			
			PH,	PL,	PT,	RO,	RU,	SC,	SD,	SE,	SG,	SK,	SL,	SY,	ΤJ,	TM,	TN,	TR,			
			TT,	TZ,	UA,	ŪĠ,	US,	UZ,	VC,	VN,	YU,	ZA,	ZM,	ZW							
		RW:	BW,	GH,	GM,	KE,	LS,	MW,	MZ,	SD,	SL,	SZ,	TZ,	UG,	ZM,	ZW,	AM,	ΑZ,			
			BY,	KG,	ΚZ,	MD,	RU,	ΤĴ,	TM,	AT,	BE,	BG,	CH,	CY,	CZ,	DE,	DK,	EE,			
•			ES,	FΙ,	FR,	GB,	GR,	HU,	ΙE,	IT,	LU,	MC,	ΝL,	PT,	RO,	SE,	SI,	SK,			
			TR,	BF,	ВJ,	CF,	CG,	CI,	CM,	GΑ,	GN,	GQ,	GW,	ML,	MR,	NE,	SN,	TD,	TG		
PRAI	KR	2002	-676	53		Α		2002	1102												
	KR 2003-75511					Α		2003	1028												

The invention discloses a composition for inhibiting the secretion of an IgE-dependent histamine-releasing factor, and pharmaceutical use thereof. The composition of the invention contains a benzimidazolic compound having proton pump inhibitor activity as an active ingredient. Furthermore, the composition of the invention may contain at least one substance selected from the group consisting of fenoctimine, oleic acid, catechin, scopadulciol, pentagalloyl glucose, bufalin, and macrolide antibiotic bafilomycin and concanamycin, all having proton pump inhibitor activity, as an addnl. active ingredient or a single active ingredient.

The composition of the invention can be advantageously used as a pharmaceutical composition for the prevention and treatment of various allergic diseases and malaria, which are caused by the IgE -dependent histamine-releasing factor.
73590-58-6, Omeprazole 102625-70-7, Pantoprazole 103577-45-3, Lansoprazole 117976-89-3, Rabeprazole RL: PAC (Pharmacological activity); THU (Therapeutic use); BIOL

(Biological study); USES (Uses)
(benzimidazolic compound with proton pump inhibitor activity for preventing secretion of IgE-dependent histamine releasing factor)

RN 73590-58-6 CAPLUS

IT

CN 1H-Benzimidazole, 5-methoxy-2-[[(4-methoxy-3,5-dimethyl-2-pyridinyl)methyl]sulfinyl]- (9CI) (CA INDEX NAME)

$$\begin{array}{c|c} N & O & Me \\ \hline N & S - CH_2 & N \\ \hline MeO & Me \\ \end{array}$$

RN 102625-70-7 CAPLUS

CN 1H-Benzimidazole, 5-(difluoromethoxy)-2-[[(3,4-dimethoxy-2-pyridinyl)methyl]sulfinyl]- (9CI) (CA INDEX NAME)

RN 103577-45-3 CAPLUS

CN 1H-Benzimidazole, 2-[[[3-methyl-4-(2,2,2-trifluoroethoxy)-2-pyridinyl]methyl]sulfinyl]- (9CI) (CA INDEX NAME)

$$\begin{array}{c|c}
N & Me \\
S - CH_2 & O - CH_2 - CF_3
\end{array}$$

RN 117976-89-3 CAPLUS

CN 1H-Benzimidazole, 2-[[[4-(3-methoxypropoxy)-3-methyl-2-pyridinyl]methyl]sulfinyl]- (9CI) (CA INDEX NAME)

$$\begin{array}{c|c}
H & O \\
N & S - CH_2
\end{array}$$
Me
$$\begin{array}{c|c}
O - (CH_2)_3 - OMe
\end{array}$$

L17 ANSWER 3 OF 4 CAPLUS COPYRIGHT 2005 ACS on STN

AN 2002:223222 CAPLUS

DN 137:257592

TI T-cell reactions to drugs in distinct clinical manifestations of drug allergy

AU Neukomm, Corinne B.; Yawalkar, Nikhil; Helbling, Arthur; Pichler, Werner J.

CS Division of Allergology, Clinic of Rheumatology and Clinical Immunology/Allergology, Inselspital, Bern, Switz.

SO Journal of Investigational Allergology and Clinical Immunology (2001), 11(4), 275-284

CODEN: JIAIEF; ISSN: 1018-9068

PB Hogrefe & Huber Publishers

DT Journal

LA English

Recent data indicate that T cells play a major role in different forms of AΒ drug allergies. To show that T-cell reactions are involved in various forms of adverse reactions to different kinds of drugs, and that lymphocyte transformation and skin tests may be pos. in patients who had distinct clin. manifestations of drug allergies. We collected data of 44 patients with a highly suggestive history for adverse drug reaction who had on subsequent investigations a pos. lymphocyte transformation test. In 41/44 patients (93%) skin tests with the suspected drugs were performed and in some cases drug-specific IgE -antibodies were determined All patients were HLA typed. Clin. manifestations of the drug allergy were heterogeneous, comprising maculopapular and bullous exanthema, erythema exsudativum multiforme, vasculitis, serum sickness, urticaria, as well as involvement of internal organs. Maculopapular exanthemas formed the largest group (54%), followed by reactions more indicative of immediate hypersensitivity (28%), such as urticaria/angioedema. In most cases (63%), β -lactam antibiotics were found to have caused the allergic reaction. Skin tests for immediate reactions were pos. in 6/40 patients (15%) tested, those for late reactions in 24/38 patients (63%) tested. Our data provide evidence that drug-specific T cells can be detected in distinct clin. manifestations of drug allergy. A combined approach using a detailed case history, lymphocyte transformation tests, skin tests (immediate and delayed type) appears to be helpful to identifying the incriminated drug.

IT 103577-45-3, Agopton

RL: ADV (Adverse effect, including toxicity); THU (Therapeutic use); BIOL (Biological study); USES (Uses)

(T-cell reactions to drugs in distinct clin. manifestations of drug allergy)

RN 103577-45-3 CAPLUS

CN 1H-Benzimidazole, 2-[[[3-methyl-4-(2,2,2-trifluoroethoxy)-2-pyridinyl]methyl]sulfinyl]- (9CI) (CA INDEX NAME)

RE.CNT 26 THERE ARE 26 CITED REFERENCES AVAILABLE FOR THIS RECORD ALL CITATIONS AVAILABLE IN THE RE FORMAT

L17 ANSWER 4 OF 4 CAPLUS COPYRIGHT 2005 ACS on STN

AN 2002:125206 CAPLUS

DN 137:210620

TI TU-572, a Potent and Selective CD45 Inhibitor, Suppresses IgE
-Mediated Anaphylaxis and Murine Contact Hypersensitivity

Reactions

ΑU Hamaguchi, Takuya; Takahashi, Akiko; Manaka, Akira; Sato, Masakazu; Osada, Hiroyuki

CS Medicinal Research Laboratories, Taisho Pharmaceutical Co., Ltd., Saitama-shi, Japan

SO International Archives of Allergy and Immunology (2001), 126(4), 318-324 CODEN: IAAIEG; ISSN: 1018-2438

S. Karger AG PΒ

DT Journal

English LA

Background: CD45, receptor-type protein tyrosine phosphatases (PTPases) AΒ are essential components of signaling through both the T cell receptor and the B cell antigen receptor. However, the functional significance of CD45 in the signaling pathway through the high-affinity Ig (Ig) E receptor has not yet been established. In this study, we demonstrate that the potent CD45 inhibitor neg. regulates IgE-dependent anaphylaxis and contact hypersensitivity reactions. Method: We have previously found that TU-572, 2-[(4-methylthiopyridin-2-yl)methylsulfinyl]-5isopropoxybenzimidazole, had a potent and selective inhibitory effect against PTPase activity of CD45. Using a CD45 inhibitor, we examined in vitro and in vivo IgE-mediated responses. Results: TU-572 potently inhibited histamine release from rat peritoneal mast cells and mouse systemic anaphylaxis reaction using monoclonal anti-dinitrophenyl (DNP) IgE and DNP-BSA. TU-572 also suppressed the immediate-type hypersensitivity response induced by repeated epicutaneous application of trinitrochlorobenzene in BALB/c mice. Conclusion: These findings revealed that the PTPase activity of CD45 played a critical role in signal transduction of IgE-mediated anaphylaxis in vitro and in vivo. PTPase inhibitors such as TU-572 are useful in the treatment of allergic diseases. IT **326592-39-6**, TU 572

RL: PAC (Pharmacological activity); THU (Therapeutic use); BIOL (Biological study); USES (Uses)

(TU-572, a potent and selective CD45 inhibitor, suppresses IgE -mediated anaphylaxis and murine contact hypersensitivity reactions)

326592-39-6 CAPLUS RN

1H-Benzimidazole, 5-(1-methylethoxy)-2-[[[4-(methylthio)-2-CN pyridinyl]methyl]sulfinyl]- (9CI) (CA INDEX NAME)

THERE ARE 37 CITED REFERENCES AVAILABLE FOR THIS RECORD 37 RE.CNT ALL CITATIONS AVAILABLE IN THE RE FORMAT

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Executing the logoff script...

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-8.76

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LOGINID:ssptamxg1614

PASSWORD:

TERMINAL (ENTER 1, 2, 3, OR ?):2

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* * * * * * * * * Welcome to STN International
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                "Ask CAS" for self-help around the clock
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NEWS 4 JAN 13 IPC 8 searching in IFIPAT, IFIUDB, and IFICDB
NEWS 5 JAN 13 New IPC 8 SEARCH, DISPLAY, and SELECT enhancements added to
                INPADOC
NEWS 6 JAN 17
                Pre-1988 INPI data added to MARPAT
NEWS 7 JAN 17 IPC 8 in the WPI family of databases including WPIFV
NEWS 8 JAN 30 Saved answer limit increased
NEWS 9 FEB 21 STN AnaVist, Version 1.1, lets you share your STN AnaVist
                visualization results
NEWS 10 FEB 22
                The IPC thesaurus added to additional patent databases on STN
NEWS 11 FEB 22 Updates in EPFULL; IPC 8 enhancements added
NEWS 12 FEB 27 New STN AnaVist pricing effective March 1, 2006
NEWS 13 FEB 28 MEDLINE/LMEDLINE reload improves functionality
NEWS 14 FEB 28 TOXCENTER reloaded with enhancements
NEWS 15 FEB 28 REGISTRY/ZREGISTRY enhanced with more experimental spectral
                property data
NEWS 16 MAR 01
                INSPEC reloaded and enhanced
NEWS 17 MAR 03 Updates in PATDPA; addition of IPC 8 data without attributes
NEWS 18 MAR 08 X.25 communication option no longer available after June 2006
NEWS 19 MAR 22 EMBASE is now updated on a daily basis
NEWS 20 APR 03 New IPC 8 fields and IPC thesaurus added to PATDPAFULL
NEWS 21 APR 03 Bibliographic data updates resume; new IPC 8 fields and IPC
                thesaurus added in PCTFULL
NEWS 22 APR 04
                STN AnaVist $500 visualization usage credit offered
                LINSPEC, learning database for INSPEC, reloaded and enhanced
NEWS 23 APR 12
NEWS 24 APR 12
                Improved structure highlighting in FQHIT and QHIT display
                in MARPAT
NEWS 25 APR 12 Derwent World Patents Index to be reloaded and enhanced during
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second quarter; strategies may be affected

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FULL ESTIMATED COST

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```
L1 ANSWER 1 OF 1 REGISTRY COPYRIGHT 2006 ACS on STN RN 326592-39-6 REGISTRY ED Entered STN: 11 Mar 2001
```

CN 1H-Benzimidazole, 5-(1-methylethoxy)-2-[[[4-(methylthio)-2-pyridinyl]methyl]sulfinyl]- (9CI) (CA INDEX NAME)

OTHER NAMES:

CN TU 572

FS 3D CONCORD

MF C17 H19 N3 O2 S2

SR CA

LC STN Files: CA, CAPLUS, CASREACT, PROUSDDR, SYNTHLINE

PROPERTY DATA AVAILABLE IN THE 'PROP' FORMAT

2 REFERENCES IN FILE CA (1907 TO DATE)
2 REFERENCES IN FILE CAPLUS (1907 TO DATE)

=> s omeprazole

L2 23 OMEPRAZOLE

=> d

L2 ANSWER 1 OF 23 REGISTRY COPYRIGHT 2006 ACS on STN

RN 755036-61-4 REGISTRY

ED Entered STN: 01 Oct 2004

CN 1H-Benzimidazole, 5-methoxy-2-[(S)-[(4-methoxy-3,5-dimethyl-2-pyridinyl)methyl]sulfinyl]-, hydrate (2:3) (9CI) (CA INDEX NAME) OTHER NAMES:

CN S-Omeprazole sesquihydrate

FS STEREOSEARCH

MF C17 H19 N3 O3 S . 3/2 H2 O

SR CA

LC STN Files: CA, CAPLUS

CRN (119141-88-7)

Absolute stereochemistry. Rotation (-).

●3/2 H₂O

1 REFERENCES IN FILE CA (1907 TO DATE)

1 REFERENCES IN FILE CAPLUS (1907 TO DATE)

=> s lansoprazole

L3 18 LANSOPRAZOLE

=> d

L3 ANSWER 1 OF 18 REGISTRY COPYRIGHT 2006 ACS on STN

RN 482370-08-1 REGISTRY

ED Entered STN: 28 Jan 2003

CN 4-Thia-1-azabicyclo[3.2.0]heptane-2-carboxylic acid, 6-[[(2R)-amino(4-hydroxyphenyl)acetyl]amino]-3,3-dimethyl-7-oxo-, (2S,5R,6R)-, mixt. with 2-[[[3-methyl-4-(2,2,2-trifluoroethoxy)-2-pyridinyl]methyl]sulfinyl]-1H-benzimidazole (9CI) (CA INDEX NAME)

OTHER NAMES:

CN Lansoprazole-amoxicillin mixt.

FS STEREOSEARCH

MF C16 H19 N3 O5 S . C16 H14 F3 N3 O2 S

CI MXS

SR CA

LC STN Files: CA, CAPLUS

CM 1

CRN 103577-45-3

CMF C16 H14 F3 N3 O2 S

$$\begin{array}{c|c}
 & \text{Me} \\
 & \text{N} \\
 & \text{N} \\
 & \text{N}
\end{array}$$

$$\begin{array}{c}
 & \text{O} \\
 & \text{CH}_2 \\
 & \text{N}
\end{array}$$

$$\begin{array}{c}
 & \text{O} \\
 & \text{CH}_2 \\
 & \text{CH}_2
\end{array}$$

CM 2

CRN 26787-78-0

CMF C16 H19 N3 O5 S

Absolute stereochemistry.

1 REFERENCES IN FILE CA (1907 TO DATE)

1 REFERENCES IN FILE CAPLUS (1907 TO DATE)

=> s pantoprazole

L4 26 PANTOPRAZOLE

=> d

L4 ANSWER 1 OF 26 REGISTRY COPYRIGHT 2006 ACS on STN

RN 867300-65-0 REGISTRY

ED Entered STN: 11 Nov 2005

CN 1H-Benzimidazol-5-ol, 6-(difluoromethoxy)-2-[[(3,4-dimethoxy-2-pyridinyl)methyl]thio]- (9CI) (CA INDEX NAME)
OTHER NAMES:

CN 6-Hydroxy-pantoprazole thioether

FS 3D CONCORD

MF C16 H15 F2 N3 O4 S

SR CA

LC STN Files: CA, CAPLUS

$$\begin{array}{c} \text{OMe} \\ \text{N} \\ \text{F}_2\text{CH} - \text{O} \end{array}$$

PROPERTY DATA AVAILABLE IN THE 'PROP' FORMAT

1 REFERENCES IN FILE CA (1907 TO DATE)

1 REFERENCES IN FILE CAPLUS (1907 TO DATE)

=> s rabeprazole

L5 6 RABEPRAZOLE

=> d

L5 ANSWER 1 OF 6 REGISTRY COPYRIGHT 2006 ACS on STN

RN 226904-99-0 REGISTRY

ED Entered STN: 01 Jul 1999

CN 1H-Benzimidazole, 2-[[[4-(3-methoxypropoxy)-3-methyl-2-

pyridinyl]methyl]sulfinyl]-, calcium salt (9CI) (CA INDEX NAME)
OTHER NAMES:

CN Rabeprazole calcium

DR 858585-57-6

MF C18 H21 N3 O3 S . 1/2 Ca

SR CA

LC STN Files: CA, CAPLUS, IMSPATENTS

CRN (117976-89-3)

●1/2 Ca

PROPERTY DATA AVAILABLE IN THE 'PROP' FORMAT

- 4 REFERENCES IN FILE CA (1907 TO DATE)
- 4 REFERENCES IN FILE CAPLUS (1907 TO DATE)

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=> LOG Y

COST IN U.S. DOLLARS

SINCE FILE

TOTAL

FULL ESTIMATED COST

ENTRY 30.74 SESSION 30.95

STN INTERNATIONAL LOGOFF AT 10:45:16 ON 14 APR 2006